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Oxytetracycline and ne- omycin sulfate amount	Indications for use	Limitations	Sponsors
	Calves (up to 250 lb): For treatment of bacterial enteritis caused by <i>E. coli</i> susceptible to oxytetracycline; treatment and control of colibacillosis (bacterial enteritis) caused by <i>E. coli</i> susceptible to neomycin	Feed continuously for 7 to 14 d; in milk replacers or starter feed. If symptoms persist after using for 2 or 3 days, consult a veterinarian. Treatment should continue 24 to 48 hours beyond remission of disease symptoms. A withdrawal period has not been established for use in preruminating calves. Do not use in calves to be processed for veal. A milk discard time has not been established for use in lactating dairy cattle. Do not use in female dairy cattle 20 months of age or older. Withdraw 5 d before slaupther.	048164 066104
	Sheep: For treatment of bacterial enteritis caused by <i>E. coli</i> and bacterial pneumonia caused by <i>P. multocida</i> susceptible to oxytetracycline; treatment and control of colibacillosis (bacterial enteritis) caused by <i>E. coli</i> susceptible to neomycin	Feed continuously for 7 to 14 d. If symptoms persist after using for 2 or 3 days, consult a veterinarian. Treatment should continue 24 to 48 hours beyond remission of disease symptoms. Withdraw 5 d before slaughter	048164 066104
(iv) To provide 25 mg/ head/day.	Calves (250 to 400 lb): For increased rate of weight gain and improved feed efficiency	Feed continuously.	048164 066104
(v) To provide 75 mg/ head/day.	Growing cattle (over 400 lb): For increased rate of weight gain; improved feed efficiency, and reduction of liver condemnation due to liver abscesses	Feed continuously.	048164 066104
(vi) To provide 0.5 to 2.0 g/head/ day.	Cattle: For prevention and treatment of the early stages of shipping fever complex	Feed 3 to 5 d before and after arrival in feedlots. A withdrawal period has not been established for use in preruminating calves. Do not use in calves to be processed for veal. A milk discard time has not been established for use in lactating dairy cattle. Do not use in female dairy cattle 20 months of age or older	048164 066104

 $[71~{\rm FR}~16225,\,{\rm Mar}.~31,\,2006,\,{\rm as}~{\rm amended}~{\rm at}~74~{\rm FR}~40724,\,{\rm Aug}.~13,\,2009]$

§558.460 Penicillin.

- (a) Specifications. As penicillin procaine G or feed grade penicillin procaine.
- (b) Sponsors. Type A medicated articles: To 066104, 100 and 227 grams per

pound. To 046573, 100 and 227 grams per

- (c) Related tolerances. See $\S 556.510$ of this chapter.
- (d) $Conditions \ of \ use.$ (1) It is used as follows:

Penicillin in grams per ton	Combination in grams per ton	Indications for use	Limitations	Sponsor
(i) 2.4 to 50		Chickens, turkeys, and pheasants; for increased rate of weight gain and improved feed efficiency	Do not feed to poultry pro- ducing eggs for human con- sumption	000069, 046573.
(ii) 5 to 20		Quail; for increased rate of weight gain and improved feed efficiency	Quail; not over 5 weeks of age.	Do.
(iii) 10 to 50		Swine; for increased rate of weight gain and improved feed efficiency		Do.

- (2) Penicillin may be used in accordance with the provisions of this section in the combinations provided as follows:
- (i) Amprolium in accordance with $\S 558.55$.
- (ii) Amprolium plus ethopatbate in accordance with $\S558.58$.

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- (iii) Hygromycin B in accordance with §558.274.
- (iv) Nicarbazin alone or with roxarsone as in §558.366.
- (v) Roxarsone and zoalene in accordance with §558.680.
- (vi) Zoalene in accordance with $\S558.680$.

[41 FR 11004, Mar. 15, 1976, as amended at 42 FR 18618, Apr. 8, 1977; 42 FR 36995, July 19, 1977; 47 FR 42103, Sept. 24, 1982; 51 FR 7399, Mar. 3, 1986; 52 FR 2686, Jan. 26, 1987; 58 FR 30120, May 26, 1993; 60 FR 39847, Aug. 4, 1995; 63 FR 36179, July 2, 1998; 65 FR 45880, July 26, 2000; 66 FR 47963, Sept. 17, 2001; 71 FR 16227, Mar. 31, 2006]

§558.464 Poloxalene.

- (a) *Approvals*. (1) Dry Type A medicated articles: 53 percent to 000069 in §510.600(c) of this chapter.
- (2) Liquid Type A medicated articles: 99.5 percent to 000069 in §510.600(c) of this chapter.
- (b) Conditions of use. (1) For prevention of legume (alfalfa, clover) and wheat pasture bloat in cattle.
- (2) Poloxalene dry Type A article and liquid Type A article must be thoroughly blended and evenly distributed in feed prior to use. This may be accomplished by adding the Type A article to a small quantity of feed, mixing thoroughly, then adding this mixture to the remaining feed and again mixing thoroughly. Dosage is 1 gram of poloxalene per 100 pounds of body weight daily and continued during exposure to bloat producing conditions. If bloating conditions are severe, the dose is doubled. Treatment should be started 2 to 3 days before exposure to bloatproducing conditions. Repeat dosage if animals are exposed to bloat-producing conditions more than 12 hours after the last treatment. Do not exceed the higher dosage levels in any 24-hour period.

[40 FR 39857, Aug. 29, 1975, as amended at 51 FR 7399, Mar. 3, 1986; 52 FR 2686, Jan. 26, 1987; 56 FR 50654, Oct. 8, 1991; 60 FR 55660, Nov. 2, 1995]

§ 558.465 Poloxalene free-choice liquid Type C feed.

- (a) *Approvals*. Type A medicated articles: 99.5 percent to 066104 in §510.600(c) of this chapter.
- (b) Conditions of use. (1) For control of legume (alfalfa, clover) and wheat pas-

ture bloat in cattle, use 7.5 grams of poloxalene per pound of liquid Type C feed (1.65 percent weight/weight). Each animal must consume 0.2 pound of Type C feed per 100 pounds of body weight daily for adequate protection.

- (2) For control of legume (alfalfa, clover) bloat in cattle grazing of prebloom legumes, use 10.00 grams of poloxalene per pound of liquid Type C feed (2.2 percent weight/weight). Each animal must consume 0.15 pound of Type C feed per 100 pounds of body weight daily for adequate protection. If consumption exceeds 0.2 pound of Type C feed per 100 pounds of body weight daily, cattle should be changed to a Type C feed containing 7.5 grams of poloxalene per pound.
- (3) Poloxalene liquid Type A article must be thoroughly blended and evenly distributed into a liquid Type C feed and offered to cattle in a covered liquid Type C feed feeder with lick wheels. The formula for the liquid Type C feed, on a weight/weight basis, is as follows: Ammonium polyphosphate 2.66 percent, phosphoric acid (75 percent) 3.37 percent, sulfuric acid 1.00 percent, water 10.00 percent, and molasses sufficient to make 100.00 percent, vitamins A and D and/or trace minerals may be added. One free-turning lick wheel per 25 head of cattle must be provided.
- (4) The medicated liquid Type C feed must be introduced at least 2 to 5 days before legume consumption to accustom the cattle to the medicated liquid Type C feed and to lick wheel feedings. If the medicated liquid wheel Type C feed feeding is interrupted, this 2- to 5-day introductory feeding should be repeated.

[40 FR 13959, Mar. 27, 1975, as amended at 42 FR 21281, Apr. 26, 1977; 51 FR 7399, Mar. 3, 1986; 52 FR 2686, Jan. 26, 1987; 56 FR 50654, Oct. 8, 1991; 60 FR 55660, Nov. 2, 1995; 66 FR 47963, Sept. 17, 2001]

§558.485 Pyrantel.

- (a) *Specifications*. Type A medicated articles containing 9.6, 19.2, 48, or 80 grams per pound pyrantel tartrate.
- (b) Approvals. See sponsors in $\S510.600$ (c) of this chapter for uses as in paragraph (e) of this section:
- (1) No. 066104: 9.6, 19.2, 48, and 80 grams per pound for use as in paragraph (e)(1) of this section.